



# SENATE MOTION

**MADAM PRESIDENT:**

**I move** that Senate Bill 406 be amended to read as follows:

- 1 Page 3, line 22, delete "of this chapter".
- 2 Page 4, after line 13, begin a new paragraph and insert:
- 3 "SECTION 6. IC 35-48-7-8.1, AS AMENDED BY P.L.131-2014,
- 4 SECTION 9, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
- 5 JULY 1, 2015]: Sec. 8.1. (a) The board shall provide for a controlled
- 6 substance prescription monitoring program that includes the following
- 7 components:
- 8 (1) Each time a controlled substance designated by the board
- 9 under IC 35-48-2-5 through IC 35-48-2-10 is dispensed, the
- 10 dispenser shall transmit to the INSPECT program the following
- 11 information:
- 12 (A) The controlled substance recipient's name.
- 13 (B) The controlled substance recipient's or the recipient
- 14 representative's identification number or the identification
- 15 number or phrase designated by the INSPECT program.
- 16 (C) The controlled substance recipient's date of birth.
- 17 (D) The national drug code number of the controlled substance
- 18 dispensed.
- 19 (E) The date the controlled substance is dispensed.
- 20 (F) The quantity of the controlled substance dispensed.
- 21 (G) The number of days of supply dispensed.
- 22 (H) The dispenser's United States Drug Enforcement Agency
- 23 registration number.
- 24 (I) The prescriber's United States Drug Enforcement Agency
- 25 registration number.
- 26 (J) An indication as to whether the prescription was
- 27 transmitted to the pharmacist orally or in writing.

- 1 (K) Other data required by the board.
- 2 (2) The information required to be transmitted under this section
- 3 must be transmitted as follows:
- 4 (A) Before July 1, 2015, not more than seven (7) days after the
- 5 date on which a controlled substance is dispensed.
- 6 (B) Beginning July 1, 2015, and until December 31, 2015, not
- 7 more than three (3) days after the date on which a controlled
- 8 substance is dispensed.
- 9 (C) Beginning January 1, 2016, and thereafter, not more than
- 10 twenty-four (24) hours after the date on which a controlled
- 11 substance is dispensed.
- 12 (3) A dispenser shall transmit the information required under this
- 13 section by:
- 14 (A) uploading to the INSPECT web site;
- 15 (B) a computer diskette; or
- 16 (C) a CD-ROM disk;
- 17 that meets specifications prescribed by the board.
- 18 (4) The board may require that prescriptions for controlled
- 19 substances be written on a one (1) part form that cannot be
- 20 duplicated. However, the board may not apply such a requirement
- 21 to prescriptions filled at a pharmacy with a Category II permit (as
- 22 described in IC 25-26-13-17) and operated by a hospital licensed
- 23 under IC 16-21, or prescriptions ordered for and dispensed to
- 24 bona fide enrolled patients in facilities licensed under IC 16-28.
- 25 The board may not require multiple copy prescription forms for
- 26 any prescriptions written. The board may not require different
- 27 prescription forms for any individual drug or group of drugs.
- 28 Prescription forms required under this subdivision must be
- 29 approved by the Indiana board of pharmacy established by
- 30 IC 25-26-13-3.
- 31 (5) The costs of the program.
- 32 **(6) Each time naloxone is dispensed, the dispenser shall**
- 33 **transmit to the INSPECT program the following information:**
- 34 **(A) The recipient's name.**
- 35 **(B) The recipient's or the recipient representative's**
- 36 **identification number or the identification number or**
- 37 **phrase designated by the INSPECT program.**
- 38 **(C) The recipient's date of birth.**
- 39 **(D) The date the naloxone is dispensed.**
- 40 **(E) The quantity of naloxone dispensed.**
- 41 **(F) The dispenser's United States Drug Enforcement**
- 42 **Agency registration number.**
- 43 **(G) An indication as to whether the prescription was**
- 44 **transmitted to the pharmacist orally or in writing.**
- 45 **(H) Other data required by the board.**
- 46 (b) This subsection applies only to a retail pharmacy. A pharmacist,

pharmacy technician, or person authorized by a pharmacist to dispense a controlled substance may not dispense a controlled substance to a person who is not personally known to the pharmacist, pharmacy technician, or person authorized by a pharmacist to dispense a controlled substance unless the person taking possession of the controlled substance provides documented proof of the person's identification to the pharmacist, pharmacy technician, or person authorized by a pharmacist to dispense a controlled substance.

SECTION 7. IC 35-48-7-10.1, AS AMENDED BY P.L.84-2010, SECTION 98, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2015]: Sec. 10.1. (a) The INSPECT program must do the following:

(1) Create a data base for information required to be transmitted under section 8.1 of this chapter in the form required under rules adopted by the board, including search capability for the following:

(A) A ~~controlled substance~~ recipient's name.

(B) A ~~controlled substance~~ recipient's or recipient representative's identification number.

(C) A ~~controlled substance~~ recipient's date of birth.

(D) The national drug code number of a controlled substance dispensed.

(E) The dates a controlled substance **or naloxone** is dispensed.

(F) The quantities of a controlled substance **or naloxone** dispensed.

(G) The number of days of supply dispensed.

(H) A dispenser's United States Drug Enforcement Agency registration number.

(I) A prescriber's United States Drug Enforcement Agency registration number.

(J) Whether a prescription was transmitted to the pharmacist orally or in writing.

(K) A controlled substance recipient's method of payment for the controlled substance **or naloxone** dispensed.

(2) Provide the board with continuing twenty-four (24) hour a day online access to the data base.

(3) Secure the information collected and the data base maintained against access by unauthorized persons.

(b) The board may execute a contract with a vendor designated by the board to perform any function associated with the administration of the INSPECT program.

(c) The INSPECT program may gather prescription data from the Medicaid retrospective drug utilization review (DUR) program established under IC 12-15-35.

1           (d) The board may accept and designate grants, public and private  
2       financial assistance, and licensure fees to provide funding for the  
3       INSPECT program."

4           Renumber all SECTIONS consecutively.  
          (Reference is to SB 406 as printed February 6, 2015.)

---

Senator MERRITT